

Amendments to the Specification

Please replace the paragraph spanning page 4, line 30- page 5, line 5, with the following replacement paragraph:

--One embodiment of the invention is a compound of Formula Ia and Ib wherein, R₁ is selected from linear, branched C₁-C₁₂-alkyl group, or cyclic C₃-C₁₂-alkyl group wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-C₆-alkyl or cyclic C₃-C₆ alkylene group or with a phenyl or phenylene group; and wherein the cyclic alkyl or cyclic alkylene group or the phenyl or phenylene group is further substituted by 0, 1, 2, or 3 methyl groups; and R₂ and R₃ are hydrogen.--

Please replace the paragraph at page 5, lines 11-15, with the following replacement paragraph:

--a) R₁ is a linear or branched C₂-C₁₁-alkyl group, or cyclic C₃-C₁₁-alkyl group wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-C₆-alkyl or cyclic C₃-C₆ alkylene group or with a phenyl or phenylene group; and wherein the cyclic alkyl or cyclic alkylene group or the phenyl or phenylene group is further substituted by 0, 1, 2, or 3 methyl groups;--

Please replace the paragraph at page 5, lines 17-21, with the following replacement paragraph:

--b) R₁ is a linear, branched or cyclic C₃-C₁₀-alkyl group, wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-C₆-alkyl or cyclic C₃-C₆ alkylene group or with a phenyl or phenylene group; and wherein the cyclic alkyl or cyclic alkylene group or the phenyl or phenylene group is further substituted by 0, 1, 2, or 3 methyl groups;--

Please replace the paragraph at page 5, lines 23-27, with the following replacement paragraph:

--c) R₁ is selected from linear, branched or cyclic C₄–C₉–alkyl group, wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-C₆-alkyl or cyclic C₃-C₆ alkylene group or with a phenyl or phenylene group; and wherein the cyclic alkyl or cyclic alkylene group or the phenyl or phenylene group is further substituted by 0, 1, 2, or 3 methyl groups; --

Please replace the paragraph spanning page 5, line 29- page 6, line 3, with the following replacement paragraph:

--d) R₁ is selected from linear, branched or cyclic C₄–C₈–alkyl group wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-C₆-alkyl or cyclic C₃-C₆ alkylene group or with a phenyl or phenylene group; and wherein the cyclic alkyl or cyclic alkylene group or the phenyl or phenylene group is further substituted by 0, 1, 2, or 3 methyl groups;--

Please replace the paragraph at page 6, lines 5-9, with the following replacement paragraph:

--e) R₁ is selected from linear, branched or cyclic C₄–C₇–alkyl group wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-C₆-alkyl or cyclic C₃-C₆ alkylene group or with a phenyl or phenylene group; and wherein the cyclic alkyl or cyclic alkylene group or the phenyl or phenylene group is further substituted by 0, 1, 2, or 3 methyl groups;--

Please replace the paragraph at page 6, lines 11-15, with the following replacement paragraph:

--f) R₁ is selected from linear, branched or cyclic C₁-C₆-alkyl group wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-C₅-alkyl or cyclic C₃-C₅ alkylene group or with a phenyl or phenylene group; and wherein the cyclic alkyl or cyclic alkylene group or the phenyl or phenylene group is further substituted by 0, 1, 2, or 3 methyl groups;--

Please replace the paragraph at page 6, lines 17-20, with the following replacement paragraph:

--g) R₁ is selected from linear, branched or cyclic C₄-alkyl group wherein the linear or branched alkyl group may be substituted or interrupted with a cyclic C₃-alkyl or cyclic C₃-alkylene group; and wherein the cyclic alkyl or cyclic alkylene group is further substituted by 0, 1, 2, or 3 methyl groups.--

Please replace the Abstract with the following replacement Abstract.

--The present invention relates to new salts of omeprazole and esomeprazole respectively, i.e. salts of 5-methoxy-2-[(4-methoxy-3,5-dimethyl-2-pyridinyl)-methyl]sulfinyl]-1*H*-benzimidazole and the (S)-enantiomer thereof. More specifically, the present invention relates to [alkylammoniumsalt] alkylammonium salts of the compounds, formed by a reaction of omeprazole and esomeprazole respectively and an alkylamine with formula [NR₂1#191R₂2#191R₂3#191] NR₁R₂R₃, wherein [R₂1#191] R₁ is a linear, branched, or cyclic [C₂1#191-C₂12#191] C₁-C₁₂-alkyl group, [R₂2#191 and R₂3#191] and R₂ and R₃ are hydrogen. The present invention also relates to a process for preparing crystalline salts, a pharmaceutical preparation, and a method for treatment of gastric related disorders by administering the compound of the invention.--